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little additive blooms, the laminated film may exhibit high COF and cause web breaks on the packaging machine's forming collar or sealing bars from excessive friction. If too much additive blooms, the laminated film may exhibit a COF that is too low for the packaging machine's tension control system, causing wrinkling or "flooding" of the line. Because of the variability in storage conditions, the same slip film using these amide systems can exhibit both extremes of COF properties.

In addition, the use of fatty amide additives also tends to make the film hazy and less glossy. This is undesirable from the point of view of print graphics appeal. Moreover, the migratory nature of the amide additive packages make them prone to bloom to the print surface of the slip film or transfer from the slip surface to the print surface when in wound roll form. The presence of amides on the print surface can interfere with the wettability and adhesion of water-based inks and, especially for process print applications, cause bridging of ink dots or inconsistent dot sizes. This results in muddier colors or images and a loss of graphic appeal.

Summary of the Invention

An objective of this invention is to solve the aforesaid problems of conventional slip films by providing an additive system that is essentially non-migratory. This will result in a slip film with stable slip and COF properties. In addition, such a film will offer excellent printability and transparency.

One aspect of the present invention is a polyolefin-based laminate film comprising at least 2 layers:

- a) a first polyolefin-based resin layer having a surface treated by a discharge treatment method that imparts excellent printability; and
- b) a polyolefin-based mixed resin layer formed on one surface of the first polyolefin-based resin layer opposite of the surface treatment,

wherein the first polyolefin-based resin layer and the polyolefin-based mixed resin layer optionally contain up to 800 ppm of fatty amides such as stearamide or erucamide and the polyolefin-based mixed resin layer contains a first additive material comprising at least one crosslinked silicone polymer in an amount of about 0.1% - 0.5% by weight of the polyolefin-based mixed resin layer and/or at least one silicone oil in an amount of about 0.02% - 0.2% by weight of the polyolefin-based mixed resin layer, and a second additive material in an amount of about 0.10 - 0.50% by weight of the polyolefin-based mixed resin layer, which comprises at least one amorphous aluminosilicate.

According to this invention, this objective and other advantages are achieved by a propylene polymer film composed of a composition comprising of at least a 2-layer coextruded film, with the print surface modified with a discharge treatment method; and the slip surface modified with a non-migratory crosslinked silicone polymer resin and/or a silicone oil and a non-migratory silicate powder additive for good slip and antiblock properties. Additionally, if amide-type additives are desired to be used, this invention allows the amount of amide additive used to be significantly reduced (less than 1000 ppm) compared to conventional slip film designs (2000 - 5000 ppm typical).

Detailed Description of the Invention

Materials that may be employed for the biaxially oriented layer are propylene homo-copolymers or copolymers of propylene and other α -olefins having 2 to 10 carbon atoms. In the case of copolymers, the amount of α -olefin subjected to copolymerization is less than 5% based on the weight of the copolymer as a standard. If the amount of copolymer exceeds this level, the biaxially oriented layer becomes too soft, with consequent insufficient service strength of the laminate film. Into the biaxially oriented layer various

additives may be introduced (normally in the range of 0.01 to 2% based on the weight of the biaxially oriented layer as a standard). These include various additives known as additives for polypropylene, for example, stabilizers, anti-oxidants, ultra-violet absorbers, plasticizers, antistatic agents, anti-blocking agents, organic lubricants, pigments, coloring agents, nucleating agents, etc. Similarly, other kinds of polymers known as suitable for mixing into polypropylene may be added, for example, polyethylene, polybutene-1, poly (4-methylpentene-1), etc. These may be added by mixing in an amount of about 0.1 to 5% based on the weight of the biaxially oriented layer.

The polyolefin-based laminate film has excellent and stable slip properties, low and stable coefficient of friction, low haze and excellent printability.

In a preferred embodiment, the first polyolefin-based resin layer has a thickness of about 6 - 40 μm . In another embodiment, this polyolefin-based resin layer is made of polypropylene-based resin. The polyolefin-based mixed resin layer has a thickness of about 0.2 - 5.0 μm .

In another embodiment, at least one component of the first additive material is a crosslinked silicone resin having a spherical average particle size of 2 - 5 μm , a specific gravity of 1.32 at 25°F, a bulk density of 0.15 - 0.50, and a linseed oil absorption rate of 50 - 90 ml/100g; and/or at least one component of the first additive material is a silicone oil having viscosity of 300 - 400 cSt., specific gravity at 77°F of 0.90 - 0.99, and volatile content of 0.001 - 0.005%.

In another embodiment, at least one component of the additive ^{is an} anti-block material ^{which} is an amorphous sodium calcium aluminosilicate having a particle size of 2 - 5 μm and a bulk density of 0.30 - 0.80 g/cm³ or an amorphous aluminosilicate having a particle size of 2 - 5 μm and a bulk density of 0.10 - 0.30 g/cm³.

5 In another embodiment, at least one component of the second additive material is an amorphous sodium calcium aluminosilicate having a particle size of 2 - 5 μm and a bulk density of 0.30 - 0.80 g/cm^3 ; or an amorphous aluminosilicate having a particle size of 2 - 5 μm and a bulk density of 0.10 - 0.30 g/cm^3 .

Example 1:

10 One hundred parts by weight of a crystalline propylene homopolymer resin; 0.0004 parts by weight of a silicone oil having a viscosity of 350 cSt. and a specific gravity of 0.97; and 0.005 parts by weight of a sodium calcium aluminosilicate powder having a mean particle diameter of 3 μm were blended together for the mixed resin layer, coextruded, and biaxially oriented to produce a 2-layer film where the mixed resin layer was 0.5 μm thick and the accompanying coextruded propylene homopolymer layer was 17 μm in thickness. The total oriented film thickness was 17.5 μm (70G or 0.7 mil thick). The film was then treated on the homopolymer side (the print surface); the mixed resin layer side was also treated but to a lesser degree (the slip side); and wound in roll form. The COF static and dynamic, hot slip properties, haze, print surface wetting tension, and printability were tested after the film was produced.

Example 2:

25 Example 1 was repeated except that 0.003 parts by weight of a crosslinked silicone polymer of 3.0 μm size were used in place of the silicone oil.

Example 3:

Example 1 was repeated except that the amount of aluminosilicate added was changed to 0.0025 parts by weight.

Example 4:

Example 2 was repeated except that the amount of aluminosilicate added was changed to 0.0025 parts by weight.

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Example 5:

Example 1 was repeated except that the amount of silicone oil was increased to 0.0008 parts by weight.

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Example 6:

Example 1 was repeated except that 0.003 parts by weight of a crosslinked silicone polymer of 3.0 μm size were added.

Example 7:

Example 6 was repeated except that the amount of aluminosilicate added was changed to 0.0025 parts by weight.

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Example 8:

Example 1 was repeated except that the accompanying coextruded propylene homopolymer layer additionally contained 0.0004 parts by weight of stearamide.

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Example 9:

Example 8 was repeated except that the stearamide was replaced with erucamide.

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Example 10:

Example 8 was repeated except that the mixed resin layer additionally contained 0.0006 parts by weight stearamide.

Comparative Example 1:

Example 1 was repeated except that no silicone oil was used.

5 Comparative Example 2:

One hundred parts by weight of a random ethylene propylene butylene
terpolymer resin; 0.003 parts by weight of erucamide fatty amide; 0.0004
parts silicone oil; and 0.003 parts by weight of a crosslinked silicone polymer
having a mean particle diameter of 4.5 μm were blended together for the
10 mixed resin layer, coextruded, and biaxially oriented to produce a 2-layer film
where the mixed resin layer was 0.5 μm thick and the accompanying
coextruded propylene homopolymer layer was 17 μm in thickness. The
accompanying coextruded propylene homopolymer layer also contained
0.0004 parts by weight steramide. The total oriented film thickness was 17.5
15 μm (70G or 0.7 mil thick). The film was then treated on the homopolymer
side (the print surface); the mixed resin layer side was also treated but to a
lesser degree (the slip side); and wound in roll form. The COF static and
dynamic, hot slip properties, haze, print surface wetting tension, and
printability were tested after the film produced.

20 Comparative Example 3:

One hundred parts by weight of a crystalline propylene homopolymer
resin; 0.003 parts by weight of behenamide and oleamide fatty amide were
blended together with 0.0025 parts by weight of an amorphous silica for the
25 mixed resin layer, coextruded, and biaxially oriented to produce a 2-layer film
where the mixed resin layer was 0.5 μm thick and the accompanying
coextruded propylene homopolymer layer was 17 μm in thickness. The total
oriented film thickness was 17.5 μm (70G or 0.7 mil thick). The film was then

treated on the homopolymer side (the print surface); the mixed resin layer side was also treated but to a lesser degree (the slip side); and wound in roll form. The COF static and dynamic, hot slip properties, haze, print surface wetting tension, and printability were tested after the film produced.

5 The various properties of the films set forth in the Examples were measured by the following methods:

A) Transparency of the film was measured by measuring the haze of a single sheet of film measured substantially in accordance with ASTM D1003. In general, the preferred value was less than 3.0% haze.

10 B) Gloss of the film was measured via a surface reflectivity gloss meter at 60°/60° angle.

C) Wetting tension of the film was measured using dyne solutions substantially in accordance with ASTM D2578.

15 D) Printability was measured by drawing down blue ink onto a single sheet of film using a meyer rod; drying the sheet in a hot air convection oven at 60°C for 1 minute; and noting the amount of ink removal when a strip of 1-inch wide 610 tape was applied and removed from the inked surface. The amount of ink removed was rated qualitatively with 5 equaling no ink removed and 1 equaling total ink removal. In general, the preferred value was 4-5.

20 E) Slipperiness of the film was measured by measuring static and kinetic COF of sheets of film via a surface property tester substantially in accordance with ASTM D1894. A 4" x 4" sled was used at ambient conditions. In general, acceptable values of the dynamic COF ranges
25 between 0.16 and 0.45; preferred values are between 0.25 and 0.35.

The properties of the films of the foregoing Examples ("Ex.") and Comparative Examples ("CEX.") are shown in Table 1.

Sample	Gauge	Haze (%)	Print Side	Print Side	Printability	Slip Side
			Gloss @ 60/60	Wetting Tension (dynes/cm)	Scale 1-5; 5=best, 1=worst	COF (st / dy)
Ex. 1	70	1.8	143.0	42	5	0.32 / 0.26
Ex. 2	70	2.2	140.3	44	5	0.28 / 0.25
Ex. 3	70	1.6	142.0	41	5	0.33 / 0.31
Ex. 4	70	2.0	141.6	41	5	0.31 / 0.28
Ex. 5	70	2.2	143.0	42	5	0.30 / 0.26
Ex. 6	70	2.7	138.0	42	5	0.23 / 0.20
Ex. 7	70	2.4	140.0	42	5	0.28 / 0.25
Ex. 8	70	2.5	141.7	42	4	0.29 / 0.25
Ex. 9	70	2.5	142.2	42	4	0.33 / 0.28
Ex. 10	70	2.6	140.1	41	4	0.32 / 0.28
CEx. 1	70	1.8	138.0	42	5	0.32 / 0.30
CEx. 2	70	2.7	129.2	41	3	0.20 / 0.18
CEx. 3	70	2.1	141.5	41	4	0.24 / 0.20

Table 1.

Table 2 shows the hot slip performance using a surface property tester modified with a 1" x 1" sled instead of a 4" x 4" sled of selected Examples and Comparative Examples. COF was measured at ambient conditions, 40°C, 60°C, 80°C, and 100°C. These conditions simulated performance under severe frictional heating that can be experienced on a packaging machine's forming collar. Test laminations of selected example films were produced for this comparison. In general, preferred hot slip performance indicated stable COF across the range of temperatures tested; at 100°C, the preference is for hot slip to be below 1.0 COF.

Sample	Hot Slip COF (st / dy)				100C
	20C	40C	60C	80C	
Ex. 1	0.33 / 0.27	0.32 / 0.28	0.30 / 0.23	0.34 / 0.24	0.55 / 0.34
Ex. 2	0.34 / 0.26	0.31 / 0.27	0.32 / 0.25	0.34 / 0.26	0.57 / 0.31
Ex. 6	0.23 / 0.20	0.25 / 0.20	0.26 / 0.21	0.33 / 0.27	0.60 / 0.36
CEx. 1	0.34 / 0.28	0.42 / 0.39	0.50 / 0.49	0.77 / 0.68	off-scale
CEx. 2	0.25 / 0.19	0.26 / 0.25	0.34 / 0.28	0.93 / 0.55	off-scale
CEx. 3	0.28 / 0.23	0.30 / 0.23	0.31 / 0.27	0.35 / 0.29	1.11 / 0.39

Table 2.

Table 3 shows the COF properties of selected samples aged for 6 months at ambient conditions in an outside warehouse under varying seasonal changes in temperature and humidity. The COF properties were compared between freshly-made film and aged film. The preferred performance should exhibit relatively stable COF performance between fresh and aged films.

Sample	COF (st / dy) "Fresh-made"	COF (st / dy) Aged 6 mos.
Ex. 1	0.32 / 0.26	0.32 / 0.29
Ex. 2	0.28 / 0.25	0.29 / 0.26
Ex. 6	0.30 / 0.26	0.31 / 0.28
CEx. 1	0.32 / 0.30	0.32 / 0.30
CEx. 2	0.19 / 0.18	0.15 / 0.13
CEx. 3	0.29 / 0.25	0.24 / 0.20

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